



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : A61K 31/56		A1	(11) International Publication Number: WO 93/15740
			(43) International Publication Date: 19 August 1993 (19.08.93)
(21) International Application Number: PCT/US93/01043 (22) International Filing Date: 29 January 1993 (29.01.93)		(81) Designated States: AU, CA, CZ, FI, JP, KR, NO, NZ, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).	
(30) Priority data: 832,828 7 February 1992 (07.02.92) US		Published <i>With international search report.</i>	
(71)(72) Applicant and Inventor: KLIGMAN, Albert, M. [US/US]; The Rittenhouse, 210 West Rittenhouse Square, Philadelphia, PA 19103 (US).			
(74) Agents: SCHWARZE, William, W. et al.; Panitch Schwarze Jacobs & Nadel, 1601 Market Street, 36th Floor, Philadelphia, PA 19103 (US).			
(54) Title: METHODS OF TREATING INFLAMMATORY DERMATOSES			
(57) Abstract			
Inflammatory dermatoses are controlled and cleared by topical application to the affected areas of the skin of a composition containing both a corticosteroid and a retinoid. The combined therapy is more effective than either active ingredient alone and is particularly effective for chronic dermatoses which are or have become resistant to corticosteroid treatment alone. After clearing has been obtained with once or twice daily applications of the corticosteroid-retinoid composition, usually after several weeks, clearance can be maintained by less frequent application or lower concentrations of the composition or by application of only one of the corticosteroid or retinoid, less potent corticosteroids, or other non-steroidal therapies, depending upon the particular dermatosis being treated.			

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FR	France	MR	Mauritania
AU	Australia	GA	Gabon	MW	Malawi
BB	Barbados	GB	United Kingdom	NL	Netherlands
BE	Belgium	GN	Guinea	NO	Norway
BF	Burkina Faso	GR	Greece	NZ	New Zealand
BG	Bulgaria	HU	Hungary	PL	Poland
BJ	Benin	IE	Ireland	PT	Portugal
BR	Brazil	IT	Italy	RO	Romania
CA	Canada	JP	Japan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SK	Slovak Republic
CI	Côte d'Ivoire	LJ	Liechtenstein	SN	Senegal
CM	Cameroon	LK	Sri Lanka	SU	Soviet Union
CS	Czechoslovakia	LU	Luxembourg	TD	Tchad
CZ	Czech Republic	MC	Monaco	TG	Togo
DE	Germany	MG	Madagascar	UA	Ukraine
DK	Denmark	ML	Mali	US	United States of America
ES	Spain	MN	Mongolia	VN	Viet Nam
FI	Finland				

- 1 -

METHODS OF TREATING INFLAMMATORY DERMATOSES

5. Field of the Invention

The present invention relates to methods for the treatment of inflammatory dermatoses. More particularly, the invention is directed to controlling, clearing and maintaining the clearance of inflammatory dermatoses by 10 administering topical compositions to the affected areas of the skin.

Background of the Invention

The most widely prescribed drugs to treat dermatologic disease are corticosteroids, also known as 15 glucocorticosteroids or glucocorticoids. Approximately 50% of prescriptions written by dermatologists are for topical corticosteroids. Since the introduction of these substances in the early 1950s for dermatologic diseases, topical corticosteroid therapy continues to be the 20 mainstay for the management of a broad spectrum of inflammatory dermatoses. Although systemic

SUBSTITUTE SHEET

- 2 -

corticosteroids are often required in some severe dermatologic diseases, topical treatment is preferred in most responsive cases because it causes fewer systemic adverse effects.

5 Topical corticosteroids are generally effective in the treatment of acute and chronic dermatoses such as seborrheic dermatitis, atopic dermatitis, contact dermatitis of the irritant and allergic type, localized neurodermatitis (lichen simplex chronicus), lichen planus,
10 and psoriasis. Steroids are also used for a variety of other less common conditions, such as Darier's disease and ichthyosiform dermatitis. A good overview of topical corticosteroid therapy appears in a series of papers presented at the Symposium on Topical Corticosteroids
15 Today and Tomorrow, sponsored by Schering AG in Bali, June 16-20, 1988, which were published in Drugs 36, Supplement 5, pp. 1-61 (Adis Press Ltd. 1988).

Individual topical corticosteroid preparations vary in anti-inflammatory potency and clinical efficacy.
20 Therapeutic efficacy of steroid therapy can often be enhanced by increasing the potency of the steroid or by using special enhancers, such as occlusive dressings. In general, efficacy is dependent on multiple factors, viz. vehicle, site and frequency of application, disease, the
25 individual patient, use of occlusive dressings, etc.

- 3 -

Potency of the corticosteroid preparation varies according to the particular corticosteroid selected, its concentration, and its vehicle. For convenience, topical corticosteroids, are classified into seven groups from 5 most (Group I) to least (Group VII) potent as shown, for example, in Table I below. Further, these classifications are ranked according to relative potency designations with Group I usually designated as ultra high potency, Groups II and III designated as high potency, Groups IV and V 10 designated mid potency, and Groups VI and VII designated low potency. Representative commercial corticosteroid preparations are set forth and classified according to this system in R. B. Stoughton, "Percutaneous Absorption of Drugs," Annual Review of Pharmacologic Toxicology, 15 pp. 55-69 (1989).

- 4 -

TABLE I
POTENCY RANKING OF TOPICAL CORTICOSTEROIDS
 (Group I, most potent to Group VII, least potent).

Group 5	Potency	Generic Name	Dosage Form	Usual Concentration
	I.	Betamethasone dipropionate	cream, ointment	0.05%
		Clobetasol propionate	cream, ointment	0.05%
		Diflorasone diacetate	ointment	0.05%
10	II.	Amcinonide	cream, ointment	0.1%
		Betamethasone dipropionate	ointment	0.05%
		Diflorasone diacetate	ointment	0.05%
		Halcinonide	cream, ointment	0.1%
		Fluocinonide	cream, ointment, solution, gel	0.05%
15		Desoximetasone	cream, gel, ointment	0.05%-0.25%
		Triamcinolone acetonide	cream, ointment	0.5%
		Mometasone	ointment	0.1%
		Fluocinolone acetonide	cream	0.2%
20	III.	Triamcinolone acetonide	ointment	0.1%
		Betamethasone dipropionate	cream	0.05%
		Diflorasone diacetate	cream	0.05%
		Betamethasone valerate	ointment	0.1%
		Mometasone	cream	0.1%
25	IV.	Flurandrenolide	ointment	0.05%
		Triamcinolone acetonide	cream, lotion	0.1%
		Fluocinolone acetonide	ointment	0.025%
		Desoximetasone	cream	0.05%
		Clocortolone pivalate	cream	0.1%
30	V.	Flurandrenolide	cream	0.05%
		Betamethasone dipropionate		0.05%
		Triamcinolone acetonide	lotion	0.1%
		Hydrocortisone butyrate	cream, ointment	0.1%
		Fluocinolone acetonide	cream	0.025%
		Betamethasone valerate	cream	0.1%
		Hydrocortisone valerate	cream	0.2%
35	VI.	Desonide	cream, ointment	0.05%
		Fluocinolone acetonide	solution	0.01
		Betamethasone valerate	lotion	0.05%
		Aciometasone dipropionate	cream, ointment	0.05%
40	VII.	Topicals with hydrocortisone, dexamethasone, flumethalone, prednisolone, and methyprednisolone		

- 5 -

Though some steroids, particularly mid- to high-potency steroids, are efficacious in chronic dermatoses, long term use of steroids is associated with serious local side effects. These include skin atrophy (thinning, telangiectasia, striae) and a prompt rebound flare when the steroid is stopped. Treatment of large areas of skin and use of occlusive dressings can also increase the potential for adverse effects. This is especially the case in children. As discussed more fully below, U.S. Patents 4,889,847 and 5,019,569 disclose the use of retinoids, such as tretinoin, to prevent and reverse skin atrophy induced by corticosteroid therapy.

Topical retinoids such as tretinoin (all-trans-retinoic acid or Vitamin A acid) have been used by dermatologists for almost twenty years. For example, tretinoin is used topically in the treatment of acne vulgaris, primarily grades I-III, in which comedones, papules, and pustules predominate. See, for example, U.S. Patent No. 3,729,568 of Kligman.

Tretinoin has been used effectively in the treatment of other skin conditions such as psoriasis, congenital ichthyosiform erythroderma, Darier's disease, epidermolytic hyperkeratosis, actinic keratosis, trichostasis, flat warts, basal cell carcinomas, and a variety of unrelated disorders. See, for example, J.R.

- 6 -

Thomas et al. "The Therapeutic Uses of Topical Vitamin A Acid," Journal of the American Academy of Dermatology, 4:505-513 (1981).

More recently, it has been found that retinoids,
5 such as tretinoin, particularly when used in separate,
sequential topical applications with the corticosteroid,
prevent and reverse skin atrophy in patients on long term
corticosteroids for various skin diseases. See, for
example, U.S. patent Nos. 4,889,847 and 5,019,569 of
10 Kligman, Mezick and Capetola, the disclosures of which are
incorporated herein by reference. However, that work was
concerned with preventing and reversing the side effects
of corticosteroid therapy and did not address the
possibility of enhanced efficacy, especially for those
15 patients whose disease has become resistant to
corticosteroids. Dermatologists call this acquired
resistance "tachyphylaxis" and try to mitigate it by
various strategies, such as rest periods (interval
therapy) and switching to another drug. These approaches
20 are only marginally helpful.

- 7 -

Brief Summary of the Invention

According to the present invention inflammatory dermatoses, including both chronic and acute varieties, can be controlled and cleared more effectively than with 5 the use of corticosteroids or retinoids alone by topically administering to the affected areas of the skin a composition comprising a corticosteroid and a retinoid in amounts which are effective for treating the dermatoses. That is, these two drugs have entirely different modes of 10 action and, when combined in a single formulation, have synergistic effects which lead to more rapid clearing and are notably effective in dermatoses which have not responded to either corticosteroids or retinoids alone. Typically, the dermatoses can be controlled and cleared by 15 once or twice daily applications of a composition containing both the retinoid and the corticosteroid in a pharmaceutically acceptable carrier for about two to three weeks.

Thereafter, clearance can be maintained by less 20 frequent and/or less potent applications of one or both of the active ingredients, such as a corticosteroid several times per week or a daily application of a retinoid. Moreover, once the disease has been brought under control, 25 lower potency steroids can be used to maintain the remission or other non-steroidal regimens can be used

- 8 -

which are safer, though usually less effective, viz. tars, topical antibiotics or anti-bacterials, and other conventional therapies. The physician is given more choices in handling inflammatory dermatoses, particularly 5 chronic inflammatory dermatoses, which are merely controlled but not cured by corticosteroids.

Detailed Description of the Preferred Embodiments

The inflammatory dermatoses which may be treated according to the present invention are well known in the 10 art. They include chronic and stubborn, as well as acute, afflictions of the skin which have previously been treated with various anti-inflammatory drugs, including oral corticosteroids and sometimes oral retinoids. While these prior therapies are sometimes effective, the side effects 15 of each are numerous and severe.

There have been reports of using tretinoin and certain corticosteroids in combination, either sequentially or mixed together, for various forms of psoriasis. See, for example, K. H. Kaidbey et al., 20 "Treatment of Psoriasis with Topically Applied Tretinoin and Steroid Ointment," Archives of Dermatology, 111:1001-1003 (1975) and P. Frost et al., "Retinoic Acid for the Therapy of Psoriasis", Acta Dermatovener, Suppl. 74:154-

- 9 -

160 (Stockholm, 1975). However, effective treatment of inflammatory dermatoses in general has not previously been indicated.

Among the disorders which can be effectively treated according to the present invention are the various forms of inflammatory acne. These include the most devastating type, acne conglobata or nodulocystic acne. Additionally, inflammatory acne with numerous pustules and deep persistent papules responds dramatically.

Severely inflammatory acne, notably acne conglobata, responds to an oral retinoid, 13-cis retinoic acid (available commercially as ACCUTANE®). However, the side effects of this drug are very serious, including teratogenicity, elevated blood lipids, fragile skin, conjunctivitis, etc. It has been found that acne conglobata as well as severely inflammatory acne vulgaris, in particular, can be brought under control or show excellent responses in as little as two to three weeks of twice daily applications of the combination treatment of the present invention. Persistent papulo-pustular acne also responds well to the combination treatment of the invention. It is noted that acne vulgaris is actually a mixture of inflammatory and non-inflammatory acnes, and while the treatments according to the present invention

- 10 -

could be used for non-inflammatory acne as well, such use would probably be unnecessary in most cases due to the effectiveness of retinoids alone.

Rosacea is another common disease which can be
5 very inflammatory and is resistant to therapy except for oral retinoids. Severe rosacea mimics acne conglobata. These fulminating types of rosacea also respond to the combination of a retinoid and a corticosteroid according to the present invention.

10 Other inflammatory disorders which may be controlled and cleared by the treatments of the present invention include lichen planus, especially the hypertrophic variety; chronic discoid lupus erythematosus; chronic atopic dermatitis, including lichen simplex
15 chronicus which is a persistent, itchy dermatosis that is common in patients with atopic dermatitis; chronic contact or allergic dermatitis, which is due to a great variety of environmental allergens; chronic hand dermatitis; lichen amyloidosis; alopecia areata; pseudofolliculitis barbae;
20 pityriasis rubra pilaris; mycosis fungoides; drug reactions (acute); and others.

The corticosteroids useful in the treatments according to the present invention include all of the large number of corticosteroids which are known for their
25 anti-inflammatory properties. See, for example, those

- 11 -

listed in Table I above. Preferably, the mid- to high- or ultra high-potency corticosteroids are used in the invention. Examples of preferred mid-potency corticosteroids are betamethasone valerate, triamcinolone 5 acetonide, and fluocinolone acetonide. Clobetasol propionate is presently unrivalled in potency. Other preferred high-potency steroids include, for example, betamethasone dipropionate. It is also possible to use low-potency corticosteroids such as hydrocortisone, 10 dexamethasone and prednisolone in those particular chronic dermatoses, such as atopic dermatitis, which do not require high-potency steroids for control.

Retinoids have been defined narrowly as comprising vitamin A (retinol) and its derivatives, such 15 as vitamin A aldehyde (retinal) and vitamin A acid (retinoic acid), which are metabolites of natural vitamin A. However, subsequent research has resulted in a larger class of chemical compounds that are termed retinoids because they have biological actions similar to 20 the parent vitamin A, even though there may be great structural dissimilarities. Compounds useful in the present invention include all natural and/or synthetic analogs of vitamin A or retinol-like compounds which have similar therapeutic activities as demonstrated for a 25 variety of retinoids. Accordingly, as used herein for

- 12 -

purposes of the present invention, the term "retinoid" will be understood to mean a natural or synthetic substance that elicits all or some of the biologic responses of retinoic acid or retinol by binding to and subsequently activating known and unknown cutaneous retinoic acid receptors. Examples of suitable retinoids useful in the present invention are set forth in Table I, although it will be understood that the invention is not limited thereto.

10

TABLE I

all-trans-retinoic acid

13-cis-retinoic acid

11-cis-retinoic acid

9-cis-retinoic acid

15

retinol

retinal

retinoyl palmitate

retinyl palmitate

retinyl propionate

20

(all-E)-9-(4-methoxy-2,3,6-

trimethylphenyl)-3,7-dimethyl-2,4,6,8-

nonatetraenoic acid ethyl ester.

- 13 -

(all-E)-9-(4-methoxy-2,3,6-trimethylphenyl)-3,7-dimethyl-2,4,6,8-nonatetraenoic acid

5 N-ethyl-9-(4-methoxy-2,3,6-trimethylphenyl)-3,7,-dimethyl-2,4,6,8-nonatetraenamide

(E,E)-9-(2,6-dichloro-4-methoxy-3-methylphenyl)-3,7-dimethyl-2,4,6,8-nonatetraenoic acid ethyl ester

10 7,8-didehydroretinoic acid

(E,E)-4-[2-methyl-4-(2,6,6-trimethyl-1-cyclohexen-1-yl)-1,3-butadienyl]benzoic acid

15 (E)-4-[4-methyl-6-(2,6,6-triemthyl-1-cyclohexen-1-yl)-1,3,5-hexatrienyl]benzoic acid

- 14 -

(all-E)-3,7-dimethyl-(3-thienyl)-
2,4,6,8-nonatetraenoic acid

(E,E,E)-3-methyl-7-(5,6,7,8-tetrahydro-
5,5,8,8y-tetramethyl-2-naphthalenyl)-
2,4,6-octatrienoic acid

5

(E)-6-[2-(2,6,6-trimethyl-1-cyclohexen-
1-yl)ethylenyl]-2-naphthalenecarboxylic acid

10

(E,E,E)-7-(2,3-dihydro-1,1,3,3-tetra-
methyl-1H-inden-5-yl)-3-methyl-
2,4,6-octatrienoic acid

(E)-4-[2-(2,3,-dihydro-1,1,3,3,-tetramethyl-
1H-inden-5-yl)-1-propenyl]benzoic acid

15

(E)-4-[2-(5,6,7,8-tetrahydro-5,5,8,8-
tetramethyl-2-naphthalenyl-1-
propenyl]benzoic acid

- 15 -

(E)-4-[2-(5,6,7,8-tetrahydro-3-methyl-
5,5,8,8-tetramethyl-2-naphthalenyl-1-
propenyl]benzoic acid

5 (E)-1,2,3,4-tetrahydro-1,1,4,4-tetramethyl-
6-(1-methyl-2-phenylethenyl)naphthalene

6-(1,2,3,4-tetrahydro-1,1,4,4-tetramethyl-
6-naphthyl-2-naphthalenecarboxylic acid

10 (E)-6-[2-(4-ethylsulfonyl)phenyl]-1-
methylethenyl]-1,2,3,4-tetrahydro-
1,1,4,4-tetramethylnaphthalene

4-[(5,6,7,8-tetrahydro-5,5,8,8-tetramethyl-
2-naphthalenyl)ethynyl]benzoic acid

15 (E)-2-(1,1,4,4-tetramethyl-1,2,3,4-tetra-
hydronaphth-7-yl)-1-[4-tetrazol-
5-yl]phenyl]-1-propene

- 16 -

(E)-4-[2-(5,6,7,8-tetrahydro-7-hydroxy-
5,5,8,8-tetramethyl-2-naphthalenyl)-
1-propenyl]benzyl alcohol

5 (6-(3-(1-adamantyl)-4-methoxyphenyl)-
2-naphthoic acid)

11-cis,13-cis-12-hydroxymethylretinoic acid δ -lactone

4-acetamidophenyl retinoate

1-(4-carboxyphenyl)-4-(5,6,7,8-tetrahydro-5,5,8,8-
tetramethyl-2-naphthalenyl)pyrazole

10 1-(4-carboxyphenyl)-5-hydroxy-3-(5,6,7,8-tetrahydro-
5,5,8,8-tetramethyl-2-naphthalenyl)pyrazole

4-[(5,6,7,8-tetrahydro-5,5,8,8-tetramethyl-2-
naphthalenyl)-ethynyl)methylsulfonylbenzene

retinoyl β -glucuronide

- 17 -

4-[(5,6,7,8-tetrahydro-5,5,8,8-tetramethyl-2-naphthalenyl)carbamoyl]benzoic acid

4-[(5,6,7,8-tetrahydro-5,5,8,8-tetramethyl-2-naphthalenyl)carboxamido]benzoic acid

5 Also encompassed within the term "retinoid" are geometric and stereoisomers of the retinoids, as well as pro-drugs thereof.

Preferably, the corticosteroid and retinoid are applied simultaneously in a single composition which uses
10 a carrier pharmaceutically acceptable for both the retinoid and corticosteroid. Since tretinoin and some other actives are relatively unstable, such as being subject to photodecomposition, the particular combination of corticosteroid, retinoid, vehicle and any other
15 ingredients must be selected so as to be compatible, but such selection is within the skill of the art given the present disclosure. The amounts or concentrations of the corticosteroid and retinoid which are present in the composition will vary widely depending upon such factors
20 as the particular corticosteroid and retinoid chosen, the disorder being treated, the frequency of applications to be made, whether or not the administration will include the use of an occlusive dressing, etc.

- 18 -

Pharmaceutical compositions containing a retinoid as an active ingredient in intimate admixture with a pharmaceutical carrier are known in the art and can be prepared according to conventional pharmaceutical compounding techniques, such as those used for formulating topical all-trans-retinoic acid (tretinoin). The carrier may take a wide variety of physical forms such as creams, dressings, gels, lotions, ointments or liquids. The particular retinoid, which may be more or less potent than tretinoin, will be present in an amount from about 0.00001% by weight to about 3% by weight, depending on the potency of the retinoid. Suitable topical retinoid preparations which are commercially available are Retin-A® gels which contain 0.01% to 0.025% by weight tretinoin, and Retin-A® creams, which contain 0.025% to 0.1% by weight tretinoin, both produced by Ortho Pharmaceutical Corporation.

Typically, the corticosteroid will be present in the composition in amounts of about 0.00001 to 3 weight percent, mainly depending on desired potency. Using tretinoin as the standard for retinoids, tretinoin will typically be present in the composition in an amount of about 0.0001 to 1 weight percent, and other more or less potent retinoids will be used in corresponding amounts equivalent thereto.

- 19 -

Low potency corticosteroids, such as hydrocortisone, will generally require concentrations of about 2% or higher, whereas mid- to high-potency corticosteroids are generally satisfactory at the 5 concentrations recommended by their manufacturers, such as 0.1% for betamethasone valerate or triamcinolone acetonide, and 0.05% for clobetasol propionate. These concentrations can of course be altered to obtain an optimal formulation for a particular condition. Moreover, 10 a particular advantage of the combinations of the present invention is that the corticosteroid can often be used in concentrations half of those in present formulations (not containing a retinoid). This is an added safety factor.

Thus, on the one hand for relatively easily 15 treated conditions, the enhanced efficacy provided by the combination with the retinoid allows the use of lower concentrations of the corticosteroid. On the other hand, for particularly resistant disorders which would not normally be controlled or cleared by either the 20 corticosteroid or retinoid alone, higher concentrations of the corticosteroid may be desired and may be more easily tolerated due to the presence of the retinoid.

The compositions of the invention may also contain additional ingredients known in the art, such as 25 stabilizers, emollients, penetration enhancers, and the

- 20 -

like. Also, the compositions may be used with various means of application including, for example, occlusive dressings and drug delivery systems such as sponges, patches, liposomes, etc.

5. The general therapeutic regimen or strategy using the combination of the present invention usually involves once or twice daily applications, preferably twice daily, of the combination for several weeks to bring the dermatoses under control. Thereafter, depending on
10. the characteristics of the disease it is possible to maintain clearance by judicious application of the corticosteroid alone several times weekly or by daily application of the retinoid alone, or by lower concentrations, and/or less frequent applications of the
15. combination or of a less potent combination.

Atopic dermatitis is an example of a disorder in which the clearance can be maintained by application of a low-potency corticosteroid alone, while inflammatory acne vulgaris is an example of a disorder in which clearance
20. can be maintained by application of a retinoid, e.g. tretinoin, alone.

Controlled studies on a significant number of patients have shown that the combined therapy according to the present invention is not only additive, but may be
25. truly synergistic. That is, the combination is more

- 21 -

effective than and produces responses not obtained by the usual treatment regimens with either the corticosteroid or the retinoid alone. Thus, inflammatory dermatoses clear more rapidly and there is rapid resolution of scaling,
5 induration and edema with the combination. This leads to greater patient compliance. Moreover, relapses are delayed and less severe. Significantly, improvement has been demonstrated in conditions which have become refractory to standard or conventional treatments, such as
10 corticosteroid therapy alone, and rebound, often found after stopping steroid therapy, is eliminated.

It is important to note that the function of the retinoid component in this combination therapy is not simply that of preventing or reversing atrophogenicity of
15 the corticosteroid, as described and claimed in U.S. Patents 4,889,847 and 5,019,569. According to the discovery of the present invention, the retinoid enhances the efficacy of the corticosteroid in suppressing inflammation which is characteristic of these chronic and
20 acute dermatoses. It is known that tretinoïn and other retinoids have some anti-inflammatory effects, though the mechanism of action is very different from that of corticosteroids. It is also known that tretinoïn makes
25 the skin more permeable, enabling a greater amount of steroid to reach the diseased skin.

- 22 -

While applicant does not wish to be bound by any particular theory of action, it is believed that the retinoid enhances the efficacy of the corticosteroid in at least two particular ways: (1) by thinning the outer 5 horny layer (the so-called permeability "barrier" of the skin), thus enabling more of the drug to penetrate into the target tissue and yielding a greater local concentration of the corticosteroid (equivalent to increasing the dose of the corticosteroid); and (2) the 10 capacity of the retinoid to prevent and reduce inflammation in its own right.

The possible explanations for the anti-inflammatory effects relate to the known ability of retinoids to inhibit migration of white cells 15 (neutrophils) from the blood vessels into the tissue (chemotaxis). Moreover, retinoids influence immune processes; for example, activation of T-cells and the release of cytokines at the site of inflammation. Retinoids also inhibit migration of macrophages into 20 diseased areas. These cells produce a variety of toxic products, including interleukins and proteins. One can cite other anti-inflammatory effects for which there is as yet no obvious explanation. Tretinoin brings about faster

- 23 -

resolution of chronic granulomas that are the result of foreign body reactions such as those elicited after the injection of collagen, elastin and carageenin.

It must be emphasized that the modes of action
5 of corticosteroids and retinoids are completely different,
and hence the combination of the two has unexpected
beneficial therapeutic effects. One mechanism by which
the corticosteroid works is by inhibiting the release of
enzymes that initiate the inflammatory cascade, whereas
10 the effect of the retinoid is less specific and dependent
on a multiplicity of unrelated effects. Among these is
the ability of retinoids to promote wound healing and to
stimulate the formation of new blood vessels
(angiogenesis), thus increasing the local blood supply.
15 Overall, retinoids, while initially inflammatory, seem to
moderate inflammatory processes. In sum, the cortico-
steroid-retinoid combinations of the invention blunt
inflammation by two entirely different mechanisms, acting
in concert.
20 Another advantage of the combination is the
prevention of rebound flare when the steroid is withdrawn.
Dramatic examples of rebound flare are well known to
dermatologists. For example, the treatment of rosacea by
topical steroids leads to a syndrome called "steroid
25 rosacea" in which atrophy and redness are prominent.

- 24 -

After cessation of treatment, a ferocious, intense, pustular eruption develops which is very difficult to control. In other diseases too, the treated sites show a "rebound dermatitis" when the steroid is stopped. The 5 site becomes tender, red, cracked, edematous and peeling. This rebound is completely prevented with combinations according to the present invention. Still further, tachyphylaxis has not been observed with the combinations according to the present invention.

10 The present invention will now be described in more detail with reference to the following specific, non-limiting examples. In these examples, two formulations have been extensively evaluated: (1) 0.1% triamcinolone acetonide combined with 0.1% tretinoin in a 15 cream base; and (2) 0.05% clobetasol propionate combined with 0.1% tretinoin in a cream base. As expected, the therapeutic response to the latter formulation was swifter, because clobetasol is the most potent corticosteroid known. Unless otherwise indicated in the 20 following examples, the combination brought about rapid resolution (control and clearing) within two to three weeks of twice daily applications.

- 25 -

Example 1

Approximately 30 cases of highly inflammatory acne vulgaris were treated with a combination of 0.05% clobetasol propionate and 0.1% tretinoin in a cream base.

- 5 Almost every instance of this condition was brought under control within two to three weeks of twice daily applications.

Example 2

- About 20 patients having severely inflammatory acne conglobata were treated with the same combination as in Example 1. These patients have shown excellent responses in three weeks of twice daily applications. The response (clearing) is faster and more dramatic than with oral 13-cis-retinoic acid.

15

Example 3

- Nine cases of severe papulo-pustular inflammatory acne were treated for two to three weeks with twice daily applications of 0.1% triamcinolone acetonide and either 0.05 or 0.1% tretinoin in a water-in-oil cream.
- 20 The lesions cooled down quickly and were then switched to daily applications of 0.025% tretinoin cream alone.

- 26 -

Another alternative after stopping the combination treatment is a topical antibiotic such as erythromycin or an anti-bacterial such as benzoyl peroxide.

Example 4

5 Approximately 14 patients with severely inflamed rosacea of the face have been treated with a combination of 0.1% tretinoin and 0.05% clobetasol propionate in a cream base. This combination was remarkably effective in clearing granulomatous rosacea, pyoderma faciale, and
10 rosacea fulminans. Thereafter, remission could be maintained by topical metronidazole or oral tetracycline.

Example 5

Six patients with hypertrophic lichen planus of the legs responded rapidly (within three weeks) from twice
15 daily applications of the same combination as in Example 4. Clearing could be maintained by mid-strength corticosteroids applied every second to third day.

Example 6

Ten adults with the adult form of chronic atopic
20 dermatitis, which appeared as lichen simplex chronicus of the lower legs were treated with a combination of 0.025% or 0.05% tretinoin and 2.0% hydrocortisone in a cream.

- 27 -

vehicle. The itchy, thick plaques which characterize this condition essentially disappeared within three weeks of twice daily applications of the combination. Thereafter, 1.0% hydrocortisone cream, applied once daily, was 5 sufficient to prevent relapse.

Example 7

A dozen patients suffering from chronic allergic contact dermatitis of occupational origin responded quickly to treatment with applications of the combination 10 of 0.1% triamcinolone acetonide and 0.1% tretinoin in a cream base.

Example 8

Twenty black males with long-standing, severely inflammatory pseudofolliculitis barbae were treated with 15 the combination of 0.1% triamcinolone acetonide and 0.1% or 0.05% tretinoin in a petrolatum vehicle (water-in-oil emulsion containing 46% petrolatum). In this disease tips of highly curved, stiff beard hairs which grow downwards into the skin create an inflammatory foreign body 20 granuloma. After about two to three weeks of treatment with the combination, the inflammatory lesions flattened and became inactive, after which tretinoin alone was used for maintenance.

- 28 -

Example 9

Small numbers of patients with various chronic dermatoses characterized by chronic, resistant, inflammatory lesions responded favorably to daily 5 applications of the combination of 0.05% clobetasol propionate and 0.1% tretinoin in a cream base for three to four weeks. These disorders included chronic discoid lupus erythematosus, lichen planus, Darier's disease, alopecia areata, and persistent seborrheic dermatitis.

10 The present invention may be embodied in other specific forms without departing from the spirit or essential attributes thereof and, accordingly, reference should be made to the appended claims, rather than to the foregoing specification as indicating the scope of the 15 invention.

- 29 -

CLAIMS

1. A method of controlling and clearing inflammatory dermatoses except psoriasis comprising topically administering to the affected area of the skin a composition comprising a corticosteroid and a retinoid, said corticosteroid and retinoid being present in amounts which are effective to control and clear said dermatosis.
2. A method according to claim 1 wherein said dermatosis is selected from the group consisting of inflammatory acne vulgaris, severe inflammatory acne conglobata, lichen planus, discoid lupus erythematosus, alopecia areata, pseudofolliculitis barbae, pityriasis rubra pilaris, mycosis fungoides, inflammatory rosacea, chronic atopic dermatitis, chronic contact dermatitis, chronic hand dermatitis, and acute drug reactions.
3. A method according to claim 1 wherein said corticosteroid is present in said composition in an amount of about 0.00001 to 3 weight percent.
4. A method according to claim 1 wherein said retinoid is present in said composition in an amount equivalent to about 0.0001 to 1 weight percent tretinoin.

- 30 -

5. A method according to claim 1 wherein said composition is applied to the skin once or twice daily until the dermatosis is controlled and cleared, and thereafter clearance is maintained by less frequent 5 application or lower concentration application of the composition or of a corticosteroid or a retinoid alone.

6. A method according to claim 5 wherein said less frequent application comprises about several applications per week of a corticosteroid or about once 10 daily applications of a retinoid.

7. A method according to claim 1 wherein said corticosteroid is a group I to group V potency corticosteroid.

8. A method according to claim 7 wherein said 15 corticosteroid is selected from the group consisting of betamethasone valerate, triamcinolone acetonide, and clobetasol propionate.

9. A method according to claim 1 wherein said retinoid is tretinoin.

- 31 -

10. A method according to claim 1 wherein said composition comprises about 0.05 to 0.1 weight percent tretinoin and about 0.05 to 0.1 weight percent of a corticosteroid selected from the group consisting of
5 betamethasone valerate, triamcinolone acetonide and clobetasol propionate.

11. A method according to claim 1 wherein said retinoid is selected from the group consisting of all-trans-retinoic acid, 13-cis-retinoic acid, 11-cis-retinoic acid, 9-cis-retinoic acid, retinol, retinal, (all-E)-9-(4-methoxy-2,3,6-trimethylphenyl)-3,7-dimethyl-2,4,6,8-nonatetraenoic acid ethyl ester, (all-E)-9-(4-methoxy-2,3,6-trimethylphenyl)-3,7-dimethyl-2,4,6,8-nonatetraenoic acid, N-ethyl-9-(4-methoxy-2,3,6-trimethylphenyl)-3,7-dimethyl-2,4,6,8-nonatetraenamide, (E,E)-9-(2,6-dichloro-4-methoxy-3-methylphenyl)-3,7-dimethyl-2,4,6,8-nonatetraenoic acid ethyl ester, 7,8-didehydroretinoic acid, (E,E)-4-[2-methyl-4-(2,6,6-trimethyl-1-cyclohexen-1-yl)-1,3-butadienyl]benzoic acid, (E)-4-[4-methyl-6-(2,6,6-trimethyl-1-cyclohexen-1-yl)-1,3,5-hexatrienyl]benzoic acid, (all-E)-3,7-dimethyl-3-thienyl)-2,4,6,8-nonatetraenoic acid, (E,E,E)-3-methyl-7-(5,6,7,8-tetrahydro-5,5,8,8-tetramethyl-2-naphthalenyl)-2,4,6-octatrienic acid, (E)-6-[2-(2,6,6-trimethyl-1-

- 32 -

cyclohexen-1-yl)ethenyl]-2-naphthalenecarboxylic acid,
(E,E,E)-7-(2,3-dihydro-1,1,3,3-tetramethyl-1H-inden-5-
yl)-3-methyl-2,4,6-octatrienoic acid, (E)-4-[2-(2,3-
dihydro-1,1,3,3-tetramethyl-1H-inden-5-yl)-1-propenyl]-
5 benzoic acid, (E)-4-[2-(5,6,7,8-tetrahydro-5,5,8,8-
tetramethyl-2-naphthalenyl-1-propenyl]benzoic acid, (E)-
4-[2-(5,6,7,8-tetrahydro-3-methyl-5,5,8,8-tetramethyl-2-
naphthalenyl-1-propenyl]benzoic acid, (E)-1,2,3,4-
tetrahydro-1,1,4,4-tetramethyl-6-(1-methyl-2-
10 phenylethenyl)-naphthalene, 6-(1,2,3,4-tetrahydro-
1,1,3,4-tetramethyl-6-naphthyl)-2-naphthalenecarboxylic
acid, (E)-6-[2-(4-(ethylsulfonyl)phenyl]-1-
methylethenyl]-1,2,3,4-tetrahydro-1,1,4,4-
tetramethylnaphthalene, 4-[(5,6,7,8-tetrahydro-5,5,8,8-
15 tetramethyl-2-naphthalenyl)ethynyl]benzoic acid, (E)-2-
(1,1,4,4-tetramethyl-1,2,3,4-tetrahydronaphth-7-yl-1-[4-
tetrazol-5-yl)phenyl]-1-propene, (E)-4-[2-(5,6,7,8-
tetrahydro-7-hydroxy-5,5,8,8-tetramethyl-2-naphthalenyl)-
1-propenyl]benzyl alcohol, retinoyl palmitate, retinyl
20 palmitate, retinyl propionate, (6-(3-(1-adamantyl)-4-
methoxyphenyl)-2-naphthoic acid), 11-cis,13-cis-12-
hydroxymethylretinoic acid β -lactone, 4-acetamidophenyl
retinate, 1-(4-carboxyphenyl)-4-(5,6,7,8-tetrahydro-
5,5,8,8-tetramethyl-2-naphthalenyl)pyrazole, 1-(4-
25 carboxyphenyl)-5-hydroxy-3-(5,6,7,8-tetrahydro-5,5,8,8-

- 33 -

tetramethyl-2-naphthalenyl)pyrazole, 4-[(5,6,7,8-tetrahydro-5,5,8,8-tetramethyl-2-naphthalenyl)-ethynyl)methylsulfonylbenzene, retinoyl β -glucuronide, 4-[(5,6,7,8-tetrahydro-5,5,8,8-tetramethyl-2-naphthalenyl)carbamoyl]benzoic acid, and 4-[(5,6,7,8-tetrahydro-5,5,8,8-tetramethyl-2-naphthalenyl)carboxamido]benzoic acid.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US92/01043

3

A. CLASSIFICATION OF SUBJECT MATTER

IPC(5) : A61K 31/56

US CL : 514/171

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/180,559

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US,A, 4,889,847 (KLIGMAN ET AL) 26 DECEMBER 1989 See entire document	1-11

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be part of particular relevance	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier document published on or after the international filing date	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&"	document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means		
"P" document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

30 APRIL 1993

Date of mailing of the international search report

26 MAY 1993

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231
Facsimile No. NOT APPLICABLEAuthorized officer *My Means*
LEONARD SCHENKMAN
Telephone No. (703) 308-1235